Acochlidium fijiensis sp. nov. (Gastropoda: Opisthobranchia: Acochlidiacea) from Fiji

by

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Abstract. A new species of freshwater opisthobranch, Acochlidium fijiensis, collected from stones in the Nasekawa River, Vanua Levu, Fiji, is described and its gross anatomical features are discussed and compared with those of other species of Acochlidium. Individual Acochlidium fijiensis reached maturity from July to October when the population was most abundant. Eggs were laid in a jelly mass attached to stones and the young hatched as veligers.

INTRODUCTION

The Acochlidiacea is the only opisthobranch order in which freshwater species are found and all of these have been discovered on islands. The freshwater species are *Strubellia paradoxa* (Strubell, 1892) from Guadalcanal, Solomon Islands, and Amboina, Indonesia (WAWRA, 1974), Acochlidi*um amboinense* Strubell, 1892, from Amboina, Acochlidium weberi Bergh, 1896, from Flores (placed in the new genus, *Palliohedyle* by RANKIN [1979] but classification disputed by Wawra [personal communication]), Acochlidium suteri Wawra, 1979, from Sumba, Indonesia, Acochlidium bayerfehlmanni Wawra, 1980, from Palau (WAWRA, 1980), and Tantulum elegans Rankin, 1979, from St. Vincent in the Caribbean (RANKIN, 1979).

MATERIALS AND METHODS

Acochlidium fijiensis was first collected 7 km upstream from the mouth of the Nasekawa River at the bridge on the Labasa-Savusavu highway on the island of Vanua Levu, the second largest island in Fiji (site, 16°40'S, 179°16'E) in October 1983 (HAYNES, 1988). Subsequently other Fijian streams and rivers have been searched for this species. The only other place where it has been found is 4 km upstream from the mouth of the Lami River, Viti Levu, the main island of Fiji (site, 18°06'S, 178°24'E). Bernadette Holthuis found 5 specimens of A. fijiensis in the Lami River in November-December 1988. Small A. fijiensis populations may exist elsewhere in Vanua Levu and Viti Levu but, because individuals are well camouflaged and blend with the stones under which they live, they are difficult to detect. Radulae were dissected from three preserved Acochlidium fijiensis. They were cleared with 10% potassium hydroxide and mounted in glycerine. The penis glands were removed from each dissected specimen, mounted in glycerine, and examined under a stereoscopic microscope.

For histology, specimens were killed and fixed in Bouin's fluid. Best results were obtained when the animals were first relaxed by lowering their temperature to about 4°C in the refrigerator. Extended, torpid specimens were immersed in ice-cold Bouin's fluid for 1 hr followed by 24 hr at room temperature. Fixed specimens were washed in running tap water for 12 hr, dehydrated in a graded series of ethanol dilutions (10–100%), cleared in xylene, and embedded in paraffin wax (melting point 60°C) under vacuum. Sections were cut at 7 μ m and stained with Erlich's haematoxylin and eosin.

Water samples were analyzed by the Institute of Natural Resources, University of the South Pacific. In general, methods for chemical analysis of water samples were those described in *Standard Methods for the Examination of Water and Wastewater* (AMERICAN PUBLIC HEALTH ASSOCIA-TION, 1981).

TAXONOMY

Acochlidium fijiensis Haynes & Kenchington,

sp. nov.

(Figures 1–6)

Type locality: Nasekawa River, Vanua Levu, Fiji. Collection site at 16°40′S, 179°16′E.

Type specimens: The holotype (LACM 2457) and 2 paratypes (LACM 2458) have been deposited in the Los







Penis gland of *Acochlidium fijiensis* showing: a, outer row of hooks; b, inner row of hooks; c, 6 small spines on the edge of the vas deferens opening.

Angeles County Museum of Natural History. Ten paratype specimens of *Acochlidium fijiensis* have been deposited in the Naturhistorisches Museum Wien, Inventory Number 84.901, and 7 paratypes, a radula, a penis permanently mounted in glycerine and slides of sectioned gonads are held in the Biology Department, School of Pure and Applied Sciences, University of the South Pacific, Suva. The dissected specimens were collected in January 1988; holotype, paratypes, and sectioned specimens were collected in July 1989. All type specimens were collected by A. Haynes.

Figure 1

Photograph of a live *Acochlidium fijiensis*. The irregular pattern of white patches over the hump and foot is caused by spicules.

Size, abundance and maturity: Table 1 indicates the relative abundance and size of specimens of Acochlidium



Figure 2A. Ventral view of a row of radula teeth of *Acochlidium fijiensis*. B. Side view of median tooth showing fine serrations.



Figure 4

Coronal (horizontal longitudinal) section of the visceral hump of *Acochlidium fijiensis* stained with haematoxylin and eosin showing the extensive distribution of ovotestis acini (A) embedded in diverticula of the digestive gland (G). Scale bar = 0.1 mm.

fijiensis collected from the Nasekawa River on five occasions from October 1983 to July 1989. Acochlidium fijiensis was most abundant in October 1983 and July 1989, and individuals were also largest in July 1989 when their body hump was enlarged because they were reproductively mature. Histological sections of A. fijiensis specimens collected from the Nasekawa River in January 1988 and from the Lami River in November-December 1988 showed undeveloped gonadal tissue. Sections of specimens collected in July 1989 contained mature gonads (Figures 4, 5). This suggests that A. fijiensis has a well-defined breeding season (July-August, or perhaps longer during the cool, dry season) and that, after breeding, either the gonads disintegrate or each individual breeds only once in its lifetime. In the mature individuals that had been prepared for histology, abundant yolk granules and sperm were noted, but very few oocytes could be seen. Presumably these specimens had already spawned.

Habitat: Specimens of *Acochlidium fijiensis* were found on the underside of stones and rocks in shallow water, 60-140 mm deep, near the water's edge in both the Nasekawa and Lami rivers. When A. fijiensis were kept in the laboratory, they always moved to the underside of the stones.

At the site in the Nasekawa River, the water level rose as much as 400 mm at high tide when heavy rain had been falling. However, when this occurred, there was little difference in the conductivity (or total ions) of the water at high and low tides (Table 2); therefore, the rise in water level is due to a back up of river water and not to inflowing seawater.

The chemical content of the water was similar at each sampling time and at high and low tides (Table 2). The water temperature was 29°C in October 1983 when Acochlidium fijiensis was abundant but at other times when the site was visited the temperature was 25-26°C (Table 2).

General Description

Diagnosis: Length of animal up to 19 mm, foot longer than visceral hump, which is rounded at the posterior (except when eggs have been shed, when the posterior may be ragged and pointed). Color cream-yellow, with wide

Figure 5

Histological sections through the ovotestis of *Acochlidium fijiensis* stained with haematoxylin and eosin. A. See abundant spermatozoa (S) and yolk granules (Y). B. See spermatozoa, yolk granules and a possible oocyte (O). Scale bar = $50 \ \mu\text{m}$.





Physical and chemical parameters at the collecting site of Acochlidium fijiensis in the Nasekawa River.

	22 October 1985		22 January 1988	
	Low tide	High tide	Low tide	High tide
Temperature (°C)	26	25	25	25
Water depth (mm)	60-140	460-600	60-140	60-160
Water speed				
$(cm \cdot s^{-1})$	0-30	0-10	0-30	0-30
pH	6.8	6.7	7.4	7.4
Conductivity				
$(\mu s \cdot cm^{-1})$	97.9	100.7	102.6	102.6
Total nitrogen				
$(mg \cdot L^{-1})$	8.2	7.1	2.5	2.5
Total phosphorus				
$(\mu g L^{-1})$	62.0	74.0	21.3	21.3
Ca (mg L^{-1})	7.8	8.2	9.2	9.2
Mg (mg L^{-1})	5.5	5.6	4.1	4.1
Na (mg L ⁻¹)	4.4	4.8	6.6	6.6
K (mg L ⁻¹)	0.26	0.29	0.85	0.85

oped penis. Also, sections of gonads (Figures 4, 5) show typical ovotestis composed of numerous acini, which are extensively distributed throughout the visceral hump where they are embedded amongst the diverticula of the digestive gland. In the specimens examined, acini were dominated by spermatozoa, spermatids, and associated generative tissue. Although yolk material was abundant, few oocytes were seen.

The spawning occurred in a 15-mm-long specimen that had been collected on 7 August 1984 and transferred to an aquarium. Thirteen days after capture it laid eggs, and three days later it died. The jelly mass was attached to two separate stones: one mass contained 31 eggs and the other 25 eggs (Figure 6A). The yellow eggs were embedded in clear jelly. After 10 days veligers were observed moving within the jelly (Figure 6B). After a further 12 days some of the veligers had escaped from the jelly mass and were swimming freely (Figure 6C). No veligers survived more than two days after leaving the jelly mass.

Discussion

A comparison of live Acochlidium fijiensis with other live Acochlidium species is not possible as no descriptions of the latter are available. Preserved specimens of A. fijiensis were similar in appearance to those described by Wawra for A. sutteri and A. bayerfehlmanni. The radula of A. fijiensis was asymmetrical, as were those of A. sutteri (WAWRA, 1979) and A. bayerfehlmanni (WAWRA, 1980). In the case of A. fijiensis, the number of rows of teeth was 50 compared with 52-56 for other Acochlidium spp. except A. weberi, which had 93-103 (WAWRA, 1979). The rachidian teeth appeared to be comparatively narrower (110 μ m compared with 200 μ m in A. amboinense and A. sutteri and 210-230 μ m in A. bayerfehlmanni) and stouter, al-



Acochlidium fijiensis. A. Developing eggs embedded in a jelly mass. B. Developing veligers, one still inside the egg membrane. C. Veliger larva swimming, after escaping from the egg membrane.

brown stripes across the dorsal side. Rhinophores (3.5 mm live; 1.8 mm preserved) longer than the anterior pair of tentacles (1.6 mm live; 0.8 mm preserved). Spicules present on visceral hump and foot (Figure 1).

Radula: Asymmetrical with formula $50 \times (1 \cdot 1 \cdot 2 \cdot)$. Left marginal plate lacking and rachidian (median) tooth finely serrated (Figure 2A, B).

Hermaphrodite: The penial armature consisting of a double row of long, curved hooks that form a border that almost surrounds the penial gland. A line of 6 small spines borders one side of the vas deferens opening (Figure 3).

Reproduction and development: Acochlidium fijiensis is a hermaphrodite. This conclusion is based on an observed spawning followed by dissection revealing a well-devel-

Table 1

The abundance and size of *Acochlidium fijiensis* collected from the Nasekawa River at different times.

Date	Number found	Duration of collection (hr)	Size (mm)
22 October 1983	16	1	not measured
7 August 1984	1	0.5	15
18-19 October 1985	12	6	6-11
19-20 January 1988	20	6	2-12
17 July 1989	30	2	10-19

(110-150 μ m) (WAWRA, 1979, 1980) (Figure 2). The male genital system of *Acochlidium fijiensis* was also similar to that of *A. sutteri* and *A. bayerfehlmanni*, with the male opening at the base of the right rhinophore (WAW-RA, 1979, 1980). The armature on the penial gland appeared to be more similar to that of *A. bayerfehlmanni* than *A. sutteri*. However, the hooks in a double row nearly surrounding the penial gland in *A. fijiensis* were long and curved and the 6 small sharp spines were in a row on one side of the penis opening (Figure 3). In *A. bayerfehlmanni* the hooks are smaller and straighter and are not so extensive.

The general anatomy of Acochlidium fijiensis more closely resembled that of A. bayerfehlmanni than A. sutteri but the penial armature was distinctly different from both. Live A. fijiensis were smaller (19 mm long) than A. bayerfehlmanni (25 mm long) (WAWRA, 1980). However, because such characteristics as the presence or absence of spicules and the comparative length of rhinophores and anterior tentacles have not previously been recorded, it is impossible to use them for comparisons within the genus Acochlidium.

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